In the claims:

1. Cancelled

2. (Currently amended) The compound according to Claim 1, or a pharmaceutically acceptable salt or stereoisomer thereof, of the Formula II:

$$R^{3a}$$
 R^{3b}
 R^{2a}
 R^{4a}
 R^{2a}
 R^{2a}

wherein

a is 0 or 1;

b is 0 or 1;

m is 0, 1 or 2;

r is 0 or 1;

s is 0 or 1;

a, b, r, s, R⁴, R⁵, R⁶ and R⁷ are defined as in Claim 1 for the compound of the Formula I and

p' is 0 to 2;

R² is selected from:

- 1) $(C=O)_aC_1-C_{10}$ alkyl,
- 2) $(C=O)_{a}$ aryl,
- 3) $(C=O)_aNR^6R^7$,
- 4) (C=O)_aC₃-C₈ cycloalkyl,
- 5) (C=O)aheterocyclyl,
- 6) SO₂NR⁶R⁷, and
- 7) SO₂C₁-C₁₀ alkyl,

said alkyl, aryl, cycloalkyl, and heterocyclyl is optionally substituted with one or more substituents selected from R⁴;

R²a is selected from: halogen and (C₁-C₆)alkyl;

R^{3a} and R^{3b} are independently selected from: hydrogen, (C₁-C₆)alkyl, trifluoromethyl and halogen; and

R^{4a} and R^{4b} are independently selected from: hydrogen, halogen and (C₁-C₆)alkyl, provided that at lease one is not hydrogen, or

R^{4a} and R^{4b} are combined to form a diradical selected from -CH₂CH₂CH₂CH₂-, -CH₂CH₂CH₂-, -CH₂CH₂-, -CH₂CH₂-, -CH₂CH₂-, -CH₂-CH₂-CH₂-, -CH₂-CH₂-CH₂-CH₂-, -CH₂-C

R4 is independently selected from:

- 1) (C=O)aObC1-C10 alkyl,
- 2) (C=O)_aO_baryl,
- 3) C2-C10 alkenyl,
- 4) C2-C10 alkynyl,
- 5) (C=O)_aO_b heterocyclyl,
- 6) CO₂H,
- 7) halo,
- 8) CN,
- 9) OH,
- 10) ObC1-C6 perfluoroalkyl,
- 11) $O_a(C=O)_bNR6R7$,
- 12) oxo,
- 13) CHO,
- 14) (N=O)R6R7,
- 15) (C=O)aObC3-C8 cycloalkyl,
- 16) SO₂NR⁶R⁷, and
- 17) SO₂C₁-C₁₀ alkyl,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one or more substituents selected from R⁵;

R⁵ is selected from:

1) $(C=O)_rO_s(C_1-C_{10})$ alkyl,

- 2) $O_r(C_1-C_3)$ perfluoroalkyl,
- 3) (C_0-C_6) alkylene-S(O)mRa,
- 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) $(C=O)_rO_s(C_2-C_{10})$ alkenyl,
- 9) $(C=O)_rO_s(C_2-C_{10})$ alkynyl,
- 10) $(C=O)_rO_s(C_3-C_6)$ cycloalkyl,
- $(C=O)_rO_s(C_0-C_6)$ alkylene-aryl,
- 12) (C=O)rOs(C0-C6)alkylene-heterocyclyl,
- 13) $(C=O)_rO_s(C_0-C_6)$ alkylene- $N(R^b)_2$,
- $C(O)R^a$,
- 15) (C₀-C₆)alkylene-CO₂R^a.
- 16) C(O)H,
- 17) (C₀-C₆)alkylene-CO₂H, and
- 18) $C(O)N(R^b)_{2}$,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b , OH, (C_1-C_6) alkoxy, halogen, CO_2H , CN, $O(C=O)C_1-C_6$ alkyl, oxo, and $N(R^b)_2$;

R6 and R7 are independently selected from:

- 1) H,
- $(C=O)O_bC_1-C_{10} \text{ alkyl},$
- 3) (C=O)ObC3-C8 cycloalkyl,
- 4) (C=O)Obaryl,
- 5) (C=O)Obheterocyclyl,
- 6) <u>C1-C10 alkyl</u>,
- <u>7)</u> aryl,
- 8) C2-C10 alkenyl,
- 9) C2-C10 alkynyl,
- 10) heterocyclyl,
- 11) C3-C8 cycloalkyl,
- 12) SO₂Ra, and
- 13) $(C=O)NRb_2$,

said alkyl, cycloalkyl, aryl, heterocylyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R⁶, or

R6 and R7 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R5;

Ra is (C1-C6)alkyl, (C3-C6)cycloalkyl, aryl, or heterocyclyl; and

Rb is H, (C1-C6)alkyl, (C1-C6)alkyl-NRa2, (C1-C6)alkyl-NH2, (C1-C6)alkyl-NHRa, aryl, heterocyclyl, (C3-C6)cycloalkyl, (C=O)OC1-C6 alkyl, (C=O)C1-C6 alkyl or S(O)2Ra.

3. (Original) The compound according to Claim 2 or a pharmaceutically acceptable salt or stereoisomer thereof, wherein:

wherein:

p' is 0 to 2; r is 0 or 1; s is 0 or 1;

R² is (C₁-C₆)alkylene-NR⁶R⁷; said alkylene is optionally substituted with up to three substituents selected from OH, (C₁-C₆)alkoxy, halogen, CO₂H, CN, O(C=O)C₁-C₆ alkyl, oxo, and NR⁶R⁷;

R²a is selected from: halogen and (C₁-C₆)alkyl;

R^{3a} and R^{3b} are independently selected from: hydrogen, (C₁-C₆)alkyl, trifluoromethyl and halogen;

R^{4a} and R^{4b} are independently selected from: hydrogen, halogen and (C₁-C₆)alkyl, provided that at least one is not hydrogen;

R⁵ is selected from:

- 1) $(C=O)_rO_s(C_1-C_{10})$ alkyl,
- 2) $O_r(C_1-C_3)$ perfluoroalkyl,
- 3) (C_0-C_6) alkylene- $S(O)_mR^a$,
- 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) $(C=O)_rO_s(C_2-C_{10})$ alkenyl,
- 9) $(C=O)_rO_s(C_2-C_{10})$ alkynyl,
- 10) $(C=O)_rO_s(C_3-C_6)$ cycloalkyl,
- 11) $(C=O)_rO_s(C_0-C_6)$ alkylene-aryl,
- 12) $(C=O)_rO_s(C_0-C_6)$ alkylene-heterocyclyl,
- 13) $(C=O)_rO_s(C_0-C_6)$ alkylene- $N(R^b)_2$,
- $C(O)R^a$,
- 15) (C₀-C₆)alkylene-CO₂R^a,
- 16) C(O)H,
- 17) (C₀-C₆)alkylene-CO₂H, and
- 18) $C(O)N(R^b)_{2}$

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b, OH, (C₁-C₆)alkoxy, halogen, CO₂H, CN, O(C=O)C₁-C₆ alkyl, oxo, and N(R^b)₂;

R6 and R7 are independently selected from:

- 1) H,
- 2) C₁-C₁₀ alkyl,
- 3) aryl,
- 4) heterocyclyl,
- 5) C2-C₁₀ alkenyl,
- 6) C2-C₁₀ alkynyl, and
- 7) C3-C8 cycloalkyl,

said alkyl, cycloalkyl, aryl, heterocylyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R⁵, or

R6 and R7 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R5;

Ra is (C1-C6)alkyl, (C3-C6)cycloalkyl, aryl, or heterocyclyl; and

Rb is H, (C1-C6)alkyl, (C1-C6)alkyl-NRa2, (C1-C6)alkyl-NH2, (C1-C6)alkyl-NHRa, aryl, heterocyclyl, (C3-C6)cycloalkyl, (C=O)OC1-C6 alkyl, (C=O)C1-C6 alkyl or S(O)₂Ra.

4. (Original) A compound which is

2-(2-bromophenyl)-3-(3-fluoro-4-methylphenyl)pyrimidin-4(3H)-one.

- 5. (Currently amended) A pharmaceutical composition that is comprised of a compound in accordance with Claim ± 2 and a pharmaceutically acceptable carrier.
- 6. (Original) A pharmaceutical composition that is comprised of a compound in accordance with Claim 3 and a pharmaceutically acceptable carrier.
 - 7. Cancelled
 - 8. Previously cancelled
 - 9. Cancelled
 - 10.- 19. Previously cancelled
 - 20.-23. Cancelled
 - 24.-26. Previously cancelled
 - 27.-33. Cancelled